Name of Committee: National Institute of Child Health and Human Development Special Emphasis Panel; Ovarian Physiology.

Date: November 1, 2021.

Time: 10:00 a.m. to 12:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Eunice Kennedy Shriver National Institute of Child Health and Human Development, National Institutes of Health, 6701 Rockledge Drive, Room 2131B, Bethesda, MD 20892 (Video Assisted Meeting).

Contact Person: Luis E. Dettin, Ph.D., Scientific Review Officer, Scientific Review Branch, Eunice Kennedy Shriver National Institute of Child Health and Human Development, National Institutes of Health, 6710B Rockledge Drive, Room 2131B, Bethesda, MD 20892, 301–827–8231, *luis.dettin@nih.gov*.

(Catalogue of Federal Domestic Assistance Program Nos. 93.865, Research for Mothers and Children, National Institutes of Health, HHS)

Dated: October 4, 2021.

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2021–21903 Filed 10–6–21; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Eunice Kennedy Shriver National Institute of Child Health and Human Development; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute of Child Health and Human Development Initial Review Group; Population Sciences Study Section.

Date: October 18, 2021.

Time: 9:00 a.m. to 2:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Eunice Kennedy Shriver National Institute of Child Health and Human Development, National Institutes of Health, 6710B Rockledge Drive, Room 2121B, Bethesda, MD 20892 (Video Assisted Meeting).

Contact Person: Christiane M. Robbins, Scientific Review Officer, Scientific Review Branch, Eunice Kennedy Shriver National Institute of Child Health and Human Development, National Institutes of Health, 6710B Rockledge Drive, Rm. 2121B, Bethesda, MD 20892, (301) 451–4989, *crobbins@mail.nih.gov.*

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

Name of Committee: National Institute of Child Health and Human Development Special Emphasis Panel; Archiving and Documenting Child Health and Human Development Data Sets.

Date: October 18, 2021.

Time: 2:00 p.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Eunice Kennedy Shriver National Institute of Child Health and Human Development, National Institutes of Health, 6710B Rockledge Drive, Room 2121B, Bethesda, MD 20892 (Video Assisted Meeting).

Contact Person: Christiane M. Robbins, Scientific Review Officer, Scientific Review Branch, Eunice Kennedy Shriver National Institute of Child Health and Human Development, National Institutes of Health, 6710B Rockledge Drive, Room 2121B, Bethesda, MD 20892, 301–451–4989, *crobbins@mail.nih.gov.*

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

(Catalogue of Federal Domestic Assistance Program Nos. 93.864, Population Research; 93.865, Research for Mothers and Children, National Institutes of Health, HHS)

Dated: October 4, 2021.

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2021–21899 Filed 10–6–21; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish a summary of information collection requests under OMB review, in compliance with the Paperwork Reduction Act. To request a copy of these documents, call the SAMHSA Reports Clearance Officer at (240) 276–0361.

Project: State Opioid Response (SOR)/ Tribal Opioid Response (TOR) Program Instrument (OMB No. 0930–0384)—Revision

SAMHSA is requesting approval to modify its existing CSAT SOR/TOR Program Instrument by (1) collapsing the original three questions into two questions for clarity and (2) adding ten questions in order to collect information on Congressionally mandated and programmatic activities, and comply with reporting requirements. The program-level information is collected quarterly and entered and stored in SAMHSA's Performance Accountability and Reporting System, which is a real-time, performance management system that captures information on the substance use prevention and treatment and mental health services delivered in the United States. Continued approval of this information collection will allow SAMHSA to continue to meet Government Performance and Results Modernization Act of 2010 reporting requirements that quantify the effects and accomplishments of its discretionary grant programs.

The SOR/TOR programs were first authorized under Title II Division H of the Consolidated Appropriations Act, 2018, Public Law 115–141. SAMHSA anticipates 159 recipients (states, territories, and tribal entities) will participate in these grant programs. Grantee-level data will include information related to: Reported overdose reversals; the purchase and distribution of naloxone; training in the administration of naloxone; implementation of prevention and education activities; outreach activities for underserved communities; and the purchase and distribution of fentanyl test strips. This grantee-level information will be collected quarterly.

The revisions to the tool will enable SAMHSA to better assess grantee accountability and performance on the required education and prevention activities for the SOR/TOR programs. SAMHSA will also use the data collected through the revised tool to implement recommendations resulting from the GAO study, ''Drug Misuse: Agencies Have Not Fully Identified How Grants That Can Support Drug Prevention Education Programs Contribute to National Goals (GAO–21–96).[1] Finally, the revisions will assist SAMHSA in providing comprehensive

[1] United States Government Accountability Office. (2020, November). Drug Misuse: Agencies Have Not Fully Identified How Grants That Can Support Drug Prevention Education Programs Contribute to National Goals. *https://www.gao.gov/assets/gao-21-96.pdf.*

data on the full range of required activities to inform Congressionally mandated reports for the SOR program.

CSAT anticipates that the time required to collect and report the program-level information is approximately 18 minutes per response. Since the submission of the original OMB package, there has been a reduction in the number of respondents. The estimated burden associated with the program-level instrument includes an adjustment to reflect the current number of grantees.

TABLE 1—ESTIMATE OF ANNUALIZED HOUR BURDEN FOR SOR/TOR GRANTEES

SAMHSA data collection	Number of respondents	Responses per respondent	Total number of responses	Burden hours per response	Total burden hours	Hourly wage [1]	Total wage cost
Grantee-Level Instrument	159	4	636	.30	190.80	$24.78	$4,728.02
CSAT Total	159	4	636	.30	190.80	24.78	4,728.02

[1] The hourly wage estimate is $24.78 based on the Occupational Employment and Wages, Mean Hourly Wage Rate for 21–1018 Substance Abuse, Behavioral Disorder, and Mental Health Counselors = $24.78/hr. as of May 2020 (*https://www.bls.gov/oes/current/oes211018.htm* Accessed on May 4, 2021.).

Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

Carlos Graham,

Reports Clearance Officer.

[FR Doc. 2021–21965 Filed 10–6–21; 8:45 am]

BILLING CODE 4162–20–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish a summary of information collection requests under OMB review, in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these documents, call the SAMHSA Reports Clearance Officer on (240) 276–0361.

Project: Opioid Drugs in Maintenance and Detoxification Treatment of Opioid Dependence—42 CFR Part 8 (OMB No. 0930–0206) and Opioid Treatment Programs (OTPs)—Extension

42 CFR part 8 establishes a certification program managed by SAMHSA's Center for Substance Abuse Treatment (CSAT). The regulation requires that Opioid Treatment Programs (OTPs) be certified. ''Certification'' is the process by which SAMHSA determines that an OTP is qualified to provide opioid treatment under the Federal opioid treatment standards established by the Secretary of Health and Human Services. To become certified, an OTP must be accredited by a SAMHSA-approved accreditation body. The regulation also provides standards for such services as individualized treatment planning, increased medical supervision, and assessment of patient outcomes. This submission seeks continued approval of the information collection requirements in the regulation and of the forms used in implementing the regulation.

SAMHSA currently has approval for the Application for Certification to Use Opioid Drugs in a Treatment Program Under 42 CFR 8.11 (Form SMA–162); the Application for Approval as Accreditation Body Under 42 CFR 8.3(b) (Form SMA–163); and the Exception Request and Record of Justification Under 42 CFR 8.12 (Form SMA–168), which may be used on a voluntary basis by physicians when there is a patient care situation in which the physician must make a treatment decision that differs from the treatment regimen required by the regulation. Form SMA–168 is a simplified, standardized form to facilitate the documentation, request, and approval process for exceptions.

SAMHSA believes that the recordkeeping requirements in the regulation are customary and usual practices within the medical and rehabilitative communities and has not calculated a response burden for them. The recordkeeping requirements set forth in 42 CFR 8.4, 8.11 and 8.12 include maintenance of the following: 5-year retention by accreditation bodies of certain records pertaining to accreditation; documentation by an OTP of the following: A patient's medical examination when admitted to treatment, a patient's history, a treatment plan, any prenatal support provided the patient, justification of unusually large initial doses, changes in a patient's dosage schedule, justification of unusually large daily doses, the rationale for decreasing a patient's clinic attendance, and documentation of physiologic dependence.

The table that follows summarizes the annual reporting burden associated with the regulation, including burden associated with the forms. There are no changes being made to the forms.

Form	Number of respondents	Responses/ respondent	Total responses	Hours/ response	Total hours
Estimated Annual Reporting Requirement Burden for Accreditation Bodies					
SMA–163	54	26.055	1,407	0.28	394
Estimated Annual Reporting Requirement Burden for Opioid Treatment Programs					
SMA–162	651.33	17.976	11,708.91		
SMA–168	1,302.67	17.977	23,418.09		
Subtotal	1954	17.977	35,127	0.08	2902
Total			36,534		3296